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Application Number

**Complete If Known**

10/500.888

Filing Date

December 1, 2004

First Named Inventor

ERICKSON et al.

### Group Art Unit

1626

**Examiner Name**

~~Thassim~~ M. Barker

Attorney Docket Number

38911-0006US1

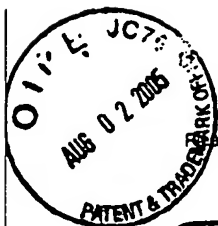
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09/13/2006

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## INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 2 of 2

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Application Number	10/500,888
Filing Date	December 1, 2004
First Named Inventor	ERICKSON et al.
Group Art Unit	1626
Examiner Name	<del>Unassigned</del> M. Barker
Attorney Docket Number	38911-0006US1

### OTHER PRIOR ART -- NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>2</sup>
MB	A02	ARUN K. GHOSH et al., "Potent HIV Protease Inhibitors Incorporating High-Affinity P2-Ligands and (R)-(Hydroxyethylamino) sulfonamide Isostere", Bioorganic & Medicinal Chemistry Letters, 1998, p. 687-690, Vol. 8	
MB	A03	ARUN K. GHOSH et al., "Structure Based Design: Novel Spirocyclic Ethers as Nonpeptidal P2-Ligands for HIV Protease Inhibitors", Bioorganic & Medicinal Chemistry Letters, 1998, pp. 979-982, Vol. 8	
MB	A04	ARUN K. GHOSH et al., "Structure-based design of non-peptide HIV protease inhibitors", IL FARMACO, January 2001, pp. 29-32, Vol. 56, No. 1/2	
MB	A05	KAZUHISA YOSHIMURA et al., "A Potent Human Immunodeficiency Virus Type 1 Protease Inhibitor, UIC-94003 (TMC-126), and Selection of a Novel (A282S) Mutation in the Protease Active Site", Journal of Virology, February 2002, pp. 1349-1358, Vol. 76, No. 3	

Examiner Signature	/Michael Barker/	Date Considered	09/13/2006
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